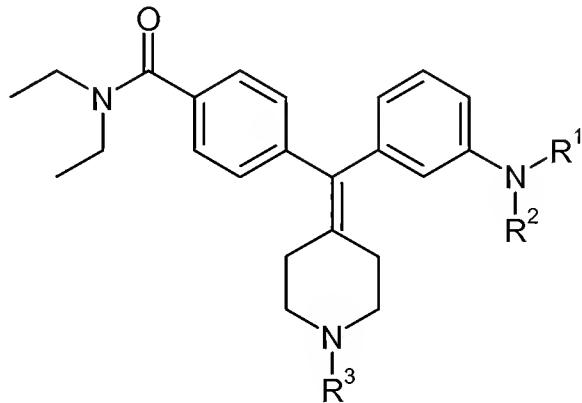


Listing of the Claims

Please amend claims 1, 2, 6-8, 10, and 18-23 as shown. The current status of all claims is listed below and supersedes all previous lists of claims.

1. (Currently amended) A compound of formula I, or a pharmaceutically acceptable salt thereof:



wherein

R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, $R^8-C(=O)-$, $R^8-S(=O)_2-$, $R^8-S(=O)-$, $R^8-NHC(=O)-$, $R^8-C(=S)-$ and $R^8-NH-C(=S)-$, wherein R^8 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl- C_{1-4} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl used in defining R^1 and R^8 are optionally substituted with one or more groups selected from -R, $-NO_2$, $-OR$, $-Cl$, $-Br$, $-I$, $-F$, $-CF_3$, $-C(=O)R$, $-C(=O)OH$, $-NH_2$, $-SH$, $-NHR$, $-NR_2$, $-SR$, $-SO_3H$, $-SO_2R$, $-S(=O)R$, $-CN$, $-OH$, $-C(=O)OR$, $-C(=O)NR_2$, $-NRC(=O)R$, and $-NRC(=O)-OR$, wherein R is, independently, selected from -H, C_{1-6} alkyl and phenyl;

R^2 is selected from -H and C_{1-6} alkyl optionally substituted with one or more groups selected from halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, and halogen, or R^1 and R^2 are $C1-3$ alkylene that together form a portion of a ring; and

R^3 is selected from -H and $-C_{1-6}$ alkyl- $O-C(=O)-$, C_{1-6} alkyl- C_{3-6} cycloalkyl, and

~~C₂₋₆cycloalkyl-C₁₋₄alkyl~~, wherein said ~~C₁₋₄alkyl-O-C(=O)-~~, C₁₋₆alkyl, ~~C₂₋₆cycloalkyl~~, and ~~C₂₋₆cycloalkyl-C₁₋₄alkyl~~ are is optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

2. (Currently amended) A compound according to claim 1, wherein

R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

R² is selected from -H and C₁₋₃alkyl; and

R³ is ~~selected from -H and C₁₋₆alkyl-O-C(=O)-~~.

3. (Original) A compound according to claim 2,

wherein R¹ is R⁹-CH₂-, wherein R⁹ is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy and halogen; and

R² and R³ are hydrogen.

4. (Original) A compound according to claim 3,

wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl, optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen.

5. (Original) A compound according to claim 4, wherein

wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl.

6. (Currently amended) A compound according to claim 1, wherein

R¹ is selected from C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

R² is -H or C₁₋₃alkyl; and

R³ is -H[[,]] or C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl are is optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen.

7. (Currently amended) A compound according to claim 6, wherein

R¹ is selected from 1-propyl, 2-propyl, 1-butyl, 2-butyl, t-butyl, 2-methyl-1-propyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, and cyclononyl;

R² is selected from -H, methyl, ethyl, 1-propyl and 2-propyl; and

R³ is selected from -H, methyl, ethyl, allyl, 3,3-dimethyl-allyl, cyclopropylmethyl, 2-methoxy-ethyl, and 3-methoxy-1-propyl.

8. (Currently amended) A compound according to claim 1, wherein

R¹ is selected from R⁸-C(=O)-, R⁸-S(=O)₂-, R⁸-S(=O)-, R⁸-NHC(=O)-, R⁸-C(=S)- and R⁸-NH-C(=S)-, wherein R⁸ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

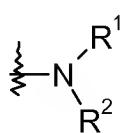
R² is -H; and

R³ is selected from -H and C₁₋₆alkyl-O-C(=O)-.

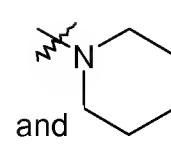
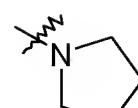
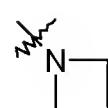
9. (Original) A compound according to claim 8, wherein

R⁸ is selected from phenyl, benzyl, phenethyl and cyclohexyl, wherein said phenyl, benzyl, phenethyl and cyclohexyl are optionally substituted with one or more groups selected from methyl, methoxy and halogen.

10. (Currently amended) A compound according to claim 1, wherein



of formula I is selected from



and

R³ is selected from -H and C₁₋₆alkyl-O-C(=O)-.

11. (Original) A compound selected from:

- 1) 4-[[3-(benzylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 2) N,N-diethyl-4-[{3-[(3-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 3) N,N-diethyl-4-(piperidin-4-ylidene){3-[(thien-3-ylmethyl)amino]phenyl}methyl)benzamide,
- 4) N,N-diethyl-4-[{3-[(2-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 5) 4-[{3-[(4-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 6) N,N-diethyl-4-[piperidin-4-ylidene(3-{[3-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,
- 7) 4-[{3-[(2-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 8) N,N-diethyl-4-[piperidin-4-ylidene(3-{[4-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,
- 9) N,N-diethyl-4-[{3-[(2-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 10) N,N-diethyl-4-(piperidin-4-ylidene){3-[(thien-2-ylmethyl)amino]phenyl}methyl)benzamide,
- 11) 4-[{3-[(cyclohexylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 12) N,N-diethyl-4-{piperidin-4-ylidene[3-(propylamino)phenyl]methyl}benzamide,
- 13) 4-[[3-(cyclohexylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 14) 4-[[3-(cyclopentylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 15) 4-[[3-(cycloheptylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 16) 4-[{3-[cyclopentyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 17) 4-[[3-(benzoylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 18) N,N-diethyl-4-[{3-[(phenylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 19) 4-[{3-[(cyclohexylcarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,

- 20) 4-[{3-[(cyclohexylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 21) 4-[(3-{{(2-chlorophenyl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 22) 4-[(3-{{(3-chlorophenyl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 23) N,N-diethyl-4-[(3-{{(5-methylthien-2-yl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 24) 4-[(3-{{(5-chlorothien-2-yl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 25) N,N-diethyl-4-[(3-{{(2S)-2-phenylpropanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 26) N,N-diethyl-4-[(3-{{(2R)-2-phenylpropanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 27) N,N-diethyl-4-[(3-{{(2S)-2-phenylbutanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 28) N,N-diethyl-4-[(3-{{(2R)-2-phenylbutanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 29) 4-[{3-[benzoyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 30) 4-[{3-[(anilinocarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 31) 4-[(3-{{(benzylamino)carbonyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 32) N-{3-[(4-[(diethylamino)carbonyl]phenyl)(piperidin-4-ylidene)methyl]phenyl}piperidine-1-carboxamide,
- 33) N,N-diethyl-4-[(3-[(phenylsulfonyl)amino]phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 34) 4-[{3-[(benzylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 35) 4-[(3-anilinophenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 36) N,N-diethyl-4-[(3-[methyl(phenyl)amino]phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 37) N,N-diethyl-4-[(3-[ethyl(phenyl)amino]phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 38) N,N-diethyl-4-[(3-{{(1S)-1-phenylethyl}amino}phenyl)(piperidin-4-

- ylidene)methyl]benzamide,
- 39) N,N-diethyl-4-[(3-{{(1R)-1-phenylethyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 40) 4-[(3-{{(1R)-1-cyclohexylethyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 41) 4-[(3-{{(1S)-1-cyclohexylethyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 42) N,N-diethyl-4-[{3-[(1-methyl-1-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 43) 4-[{3-[cyclohexyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 44) N,N-diethyl-4-[piperidin-4-ylidene(3-piperidin-1-ylphenyl)methyl]benzamide,
- 45) N,N-diethyl-4-[piperidin-4-ylidene(3-pyrrolidin-1-ylphenyl)methyl]benzamide,
- 46) N,N-diethyl-4-[[3-[(2-ethyl-1-oxobutyl)amino]phenyl]-4-piperidinyldenemethyl]-benzamide,
- 47) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-1-methyl-1H-1,2,3-benzotriazole-5-carboxamide,
- 48) 6-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-3-pyridinecarboxamide,
- 49) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-2-methoxybenzamide,
- 50) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-2-quinoxalinecarboxamide,
- 51) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-2,5-difluorobenzamide,
- 52) 3-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-2-thiophenecarboxamide,
- 53) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinyldenemethyl]phenyl]-3-methylbenzamide,
- 54) N,N-diethyl-4-[[3-[(methylphenylamino)carbonyl]amino]phenyl]-4-piperidinyldenemethyl]-benzamide, and pharmaceutically acceptable salts thereof.

12-13. (Cancelled)

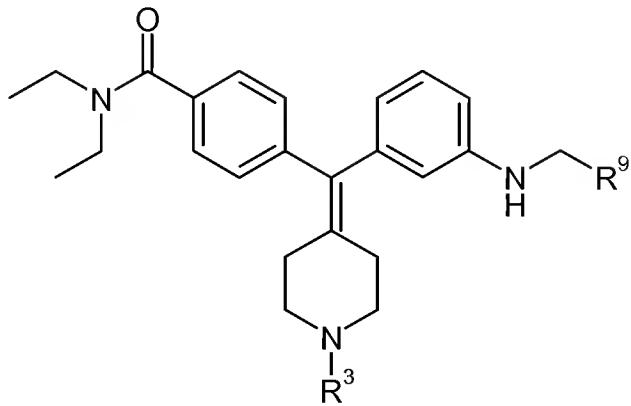
14. (Withdrawn) A pharmaceutical composition comprising a compound according to claim 1 and a pharmaceutically acceptable carrier.

15. (Withdrawn) A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

16. (Withdrawn) A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

17. (Withdrawn) A method for the therapy of anxiety in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

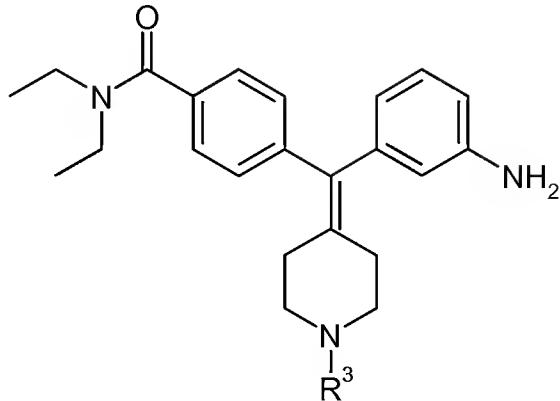
18. (Currently amended) A process for preparing a compound of formula III,



III

comprising:

reacting a compound of formula II,

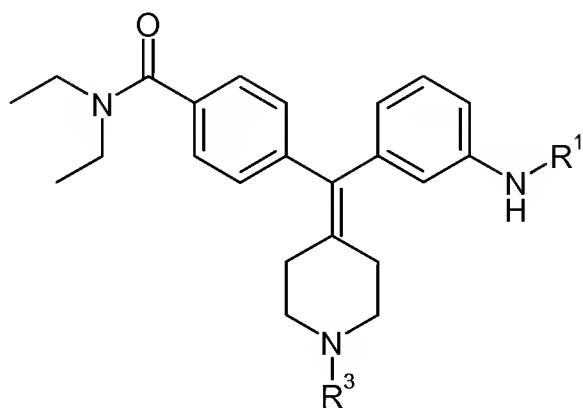


with $R^9\text{-CHO}$ in the presence of a reducing agent to form the compound of formula III,
wherein

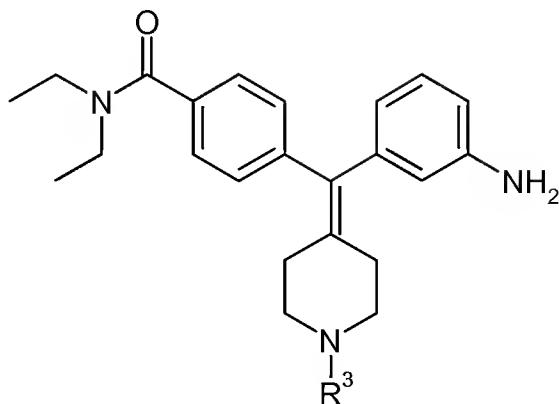
R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from $C_{1-4}\text{alkyl}$, halogen, $-\text{CF}_3$, $-\text{OH}$, $C_{1-3}\text{alkoxy}$, phenoxy and halogen; and

R^3 is selected from $C_{1-6}\text{alkyl-O-C(=O)}$, $C_{1-6}\text{alkyl}$, $C_{2-6}\text{cycloalkyl}$, and $C_{2-6}\text{cycloalkyl-C}_{1-4}\text{alkyl}$, wherein said $C_{1-6}\text{alkyl-O-C(=O)}$, $C_{1-6}\text{alkyl}$, $C_{2-6}\text{cycloalkyl}$, and $C_{2-6}\text{cycloalkyl-C}_{1-4}\text{alkyl}$ are which is optionally substituted with one or more groups selected from $C_{1-6}\text{alkyl}$, halogenated $C_{1-6}\text{alkyl}$, $-\text{NO}_2$, $-\text{CF}_3$, $C_{1-6}\text{alkoxy}$ and halogen.

19. (Currently amended) A process for preparing a compound of formula IV,



comprising: reacting a compound of formula II,

**II**

with R¹-X to form the compound of formula IV,

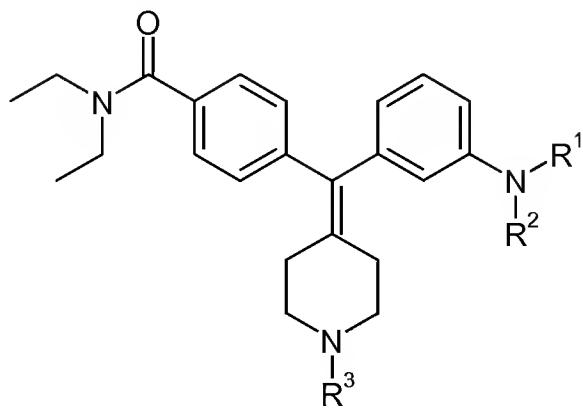
wherein

X is halogen;

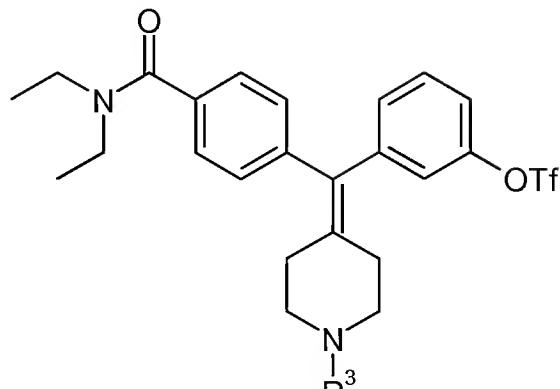
R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

~~R³ is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl-, C₂₋₆cycloalkyl, and C₂₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl-, C₂₋₆cycloalkyl, and C₂₋₆cycloalkyl-C₁₋₄alkyl are which is~~ optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

20. (Currently amended) A process for preparing a compound of formula I,

**I**

comprising: reacting a compound of formula V,



V

with R¹R²NH to form the compound of formula I,

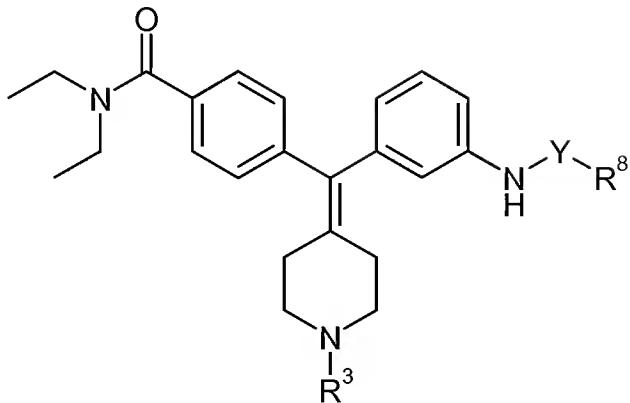
wherein

R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

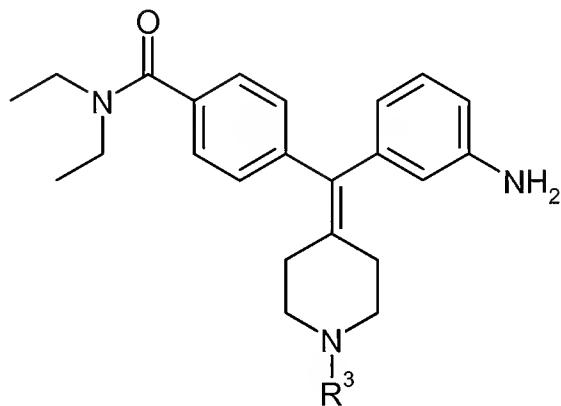
R² is selected from -H and C₁₋₆alkyl optionally substituted with one or more groups selected from halogen, -CF₃, -OH, C₁₋₃alkoxy, and halogen, or R¹ and R² are C₁₋₃alkylene that together form a portion of a ring; and

R³ is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are which is optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

21. (Currently amended) A process for preparing a compound of formula VI,



comprising: reacting a compound of formula VII,



with R⁸-Y-X or R⁸-Y-O-Y-R⁸ to form the compound of formula VI:

wherein

X is halogen;

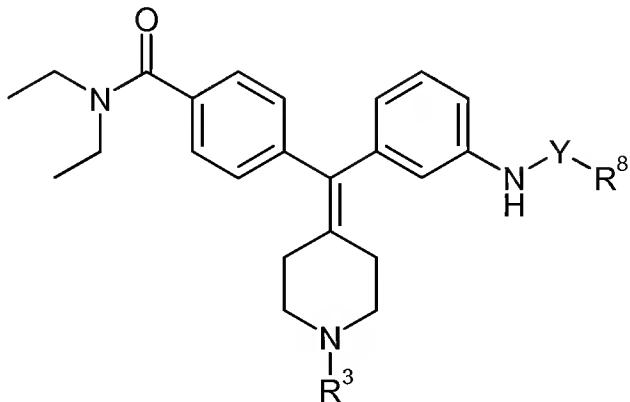
Y is selected from -C(=O)- and -S(=O)₂-;

R⁸ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

R³ is selected from C₄₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₄₋₆alkyl, wherein said C₄₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₄₋₆alkyl are which is optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated

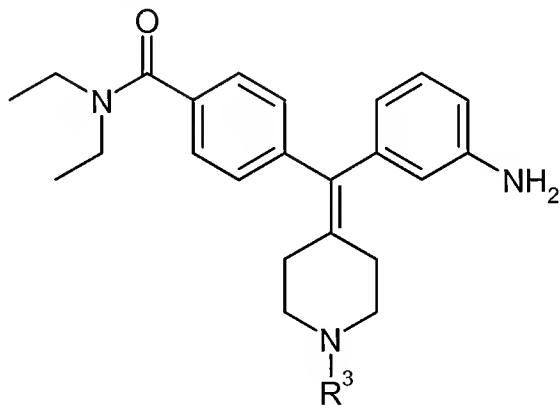
C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

22. (Currently amended) A process for preparing a compound of formula VIII,



VIII

comprising: reacting a compound of formula VII,



VII

with R⁸-Z to form the compound of formula VIII:

wherein

Z is selected from -NCO and -NCS;

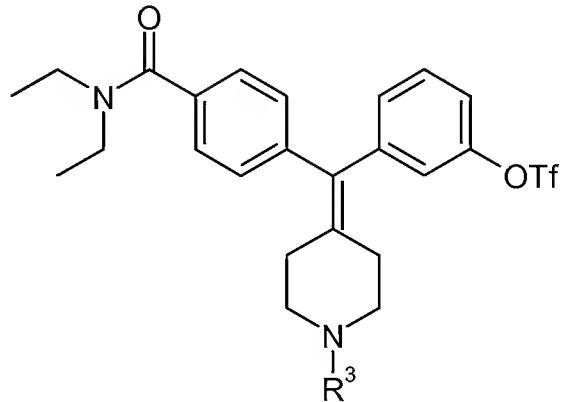
Y is selected from -C(=O)NH- and -C(=S)NH-;

R⁸ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

R³ is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄

~~alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₂₋₆cycloalkyl, and C₂₋₆cycloalkyl-C₁₋₄alkyl are which is~~ optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

23. (Currently amended) A compound of formula V,



V

wherein

R³ is selected from ~~C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₂₋₆cycloalkyl, and C₂₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₂₋₆cycloalkyl, and C₂₋₆cycloalkyl-C₁₋₄alkyl are which is~~ optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.